## **AMENDMENTS TO THE CLAIMS**

This listing of claims will replace all prior versions and listings of claims in the application:

## LISTING OF CLAIMS:

1. (Currently amended) A method for preparing a thieno[3,2-c]pyridine derivative of formula (1) comprising reacting a compound of formula (2e) with a compound of formula (3) or its salt:

$$R$$
 $S$ 
 $Y'$ 
 $(2e)$ 
 $R$ 
 $H_2N$ 
 $(3)$ 

wherein,

R is hydrogen or methoxycarbonyl; and

X' and Y' are each independently chloro, bromo, methanesulfonylmethanesulfonyloxy or  $\underline{p\text{-toluenesul} fonyl} \underline{p\text{-toluenesul} fonyloxy}.$ 

2. (Original) The method of claim 1, wherein the compound of formula (2e) is obtained by (a) cyclizing a compound of formula (4) with 2,5-dihydroxy-1,4-dithiane to obtain a compound of formula (2a), (b) reducing the compound of formula (2a) with a reducing agent to obtain a compound of formula (2b), and (c) reacting the compound of formula (2b) with a halogenating or sulfonylating agent:

$$R_6O_2C$$
  $CO_2R_7$   $(4)$ 

wherein,

R<sub>3</sub> and R<sub>4</sub> are each independently hydrogen or straight or branched C<sub>1-6</sub> alkyl, and  $R_6$  and  $R_7$  are each independently straight or branched  $C_{1\text{-}6}$  alkyl.

3. (Original) The method of claim 1, wherein the compound of formula (2e) is obtained by (a) cyclizing directly 2-thiopheneethanol with formylating agent, or reacting 2-thiopheneethanol with dialkoxymethane to obtain a compound of formula (2c) and then cyclizing the compound of formula (2c), to obtain the compound of formula (2d) and (b) reacting the compound of formula (2d) with a halogenating agent:

wherein,

 $R_5$  is  $C_{1-4}$  alkoxymethyl.

- 4. (Currently Amended) The method of claim 1, wherein the compound of formula (3) is 2-chlorobenzylamine or (S) (+)-2-chlorophenylglycin(S)-(+)-2-(2-chlorophenyl)glycine methyl ester, or a salt thereof.
- 5. (Original) The method of claim 1, wherein the compound of formula (3) or its salt is employed in an amount of 1 to 2 molar equivalents based on the amount of the compound of formula (2e).

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6. (Original) The method of claim 1, wherein the reaction is conducted in an organic solvent in the presence of a base.

- 7. (Original) The method of claim 6, wherein the organic solvent is selected from the group consisting of tertiary alcohols, ethers, nitriles, esters, optionally halogenated hydrocarbons, amides, toluene, dimethylsulfoxide and a mixture thereof.
- 8. (Original) The method of claim 6, wherein the base is an organic base selected from the group consisting of triethylamine, diisopropylethylamine, tributylamine, pyridine, picoline and a mixture thereof, or an inorganic base selected from the group consisting of sodium hydrogen carbonate, sodium carbonate, potassium hydrogen carbonate, potassium carbonate, sodium hydrogen phosphate, potassium hydrogen phosphate and a mixture thereof, or a combination thereof.
- 9. (Original) The method of claim 6, wherein the base is employed in an amount of 2 to 5 molar equivalents based on the amount of the compound of formula (2e).
- 10. (Currently Amended) The method of claim <u>6</u>1, wherein the reaction is carried out at a temperature ranging from room temperature to the boiling point of the solvent used.

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11. (Withdrawn) A compound of formula (2a) as an intermediate for the preparation of a thieno[3,2-c]pyridine derivative of formula (1) according to claim 1:

$$CO_2R_3$$
 $CO_2R_4$ 
 $(2a)$ 

wherein,

R<sub>3</sub> and R<sub>4</sub> are each independently hydrogen or straight or branched C<sub>1-6</sub> alkyl.

12. (Withdrawn) A compound of formula (2b) as an intermediate for the preparation of a thieno[3,2-c]pyridine derivative of formula (1) according to claim 1.

13. (Withdrawn) A compound of formula (2c) as an intermediate for the preparation of a thieno[3,2-c]pyridine derivative of formula (1) according to claim 1:

wherein,

 $R_5$  is  $C_{1-4}$  alkoxymethyl.

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14. (Withdrawn) A compound of formula (2d) as an intermediate for the preparation

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15. (Currently Amended) A compound of formula (2e) as an intermediate for the preparation of a thieno[3,2-c]pyridine derivative of formula (1) according to claim 1:

of a thieno[3,2-c]pyridine derivative of formula (1) according to claim 1.

wherein,

 $X' \ and \ Y' \ are \ each \ independently \ chloro, \ bromo, \ \frac{methanesulfonyl methanesulfonyloxy}{methanesulfonyloxy} \ or \\ \frac{p-toluenesulfonyl p-toluenesulfonyloxy}{methanesulfonyloxy}.$